Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

Claim 4 (previously presented): The method of claim 15 wherein the contacting is performed *in vitro*.

Claim 5 (previously presented): The method of claim 15 wherein the contacting is performed *in vivo*.

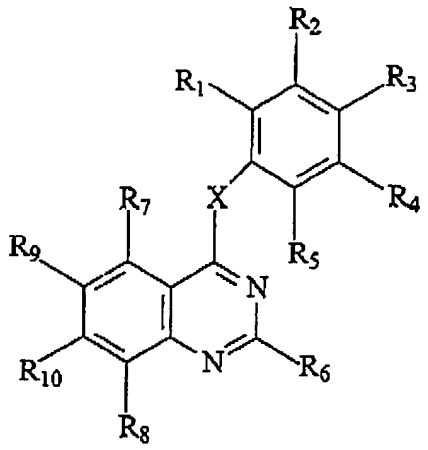
Claim 10 (currently amended): The method of claim 15 wherein the compound is 4-(4'-hydroxylphenyl)-amino-6,7-dimethoxyquinazoline ~~or~~ ~~4-(3'-bromo-4'-hydroxylphenyl)-amino-6,7-dimethoxyquinazoline;~~ or a pharmaceutically acceptable salt thereof.

Claim 11 (previously presented): The method of claim 15 wherein the cells are mammalian.

Claim 12 (previously presented): The method of claim 15 wherein the cells are human.

Claim 13 (previously presented): The method of claim 15 wherein the cells are avian.

Claim 15 (currently amended): A method for specifically inhibiting c-jun activation in mammalian or avian cells comprising contacting the cells with an effective inhibitory amount of a compound of formula I:



wherein

X is HN, $R_{11}N$, S, O, CH_2, or $R_{11}CH$;

R_{11} is hydrogen (C_1-C_4)alkyl, or (C_1-C_4)alkanoyl;

R_1-R_8 are each independently hydrogen, hydroxy, mercapto, amino, nitro, (C_1-C_4)alkyl, (C_1-C_4)alkoxy, (C_1-C_4)alkylthio, or halo; wherein two adjacent groups of R_1-R_5 together with the phenyl ring to which they are attached may optionally form a fused ring~~, for example forming a naphthyl or a tetrahydronaphthyl ring~~; and further wherein the ring formed by the two adjacent groups of R_1-R_5 may optionally be substituted by 1, 2, 3, or 4 hydroxy, mercapto, amino, nitro, (C_1-C_4)alkyl, (C_1-C_4)alkoxy, (C_1-C_4)alkylthio, or halo; and

R_9 and R_{10} are each independently hydrogen, (C_1-C_4)alkyl, (C_1-C_4)alkoxy, halo, or (C_1-C_4)alkanoyl; or R_9 and R_{10} together are methylenedioxy; or a pharmaceutically acceptable salt thereof.

Claim 16 (new): The method of claim 15 wherein the compound is

4-(3'-bromo-4'-hydroxylphenyl)-amino-6,7-dimethoxyquinazoline or

a pharmaceutically acceptable salt thereof.